



## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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<b>(21) International Application Number:</b> PCT/US00/09798  <b>(22) International Filing Date:</b> 12 April 2000 (12.04.00)  <b>(30) Priority Data:</b> 60/129,610                      16 April 1999 (16.04.99)                      US  <b>(71) Applicant:</b> SCHERING CORPORATION [US/US]; 2000 Galloping Hill Road, Kenilworth, NJ 07033 (US).  <b>(72) Inventors:</b> ALTMANN, Scott, W.; 42 First Street, Fanwood, NJ 07023 (US). BURNETT, Duane, A.; 9 Chestnut Avenue, Bernardsville, NJ 07924 (US). DAVIS, Harry, R., Jr.; 64 River Bend Road, Berkeley Heights, NJ 07922 (US). GRAZIANO, Michael, P.; 2239 Concord Road, Scotch Plains, NJ 07076 (US). LAVERTY, Maureen; 36 Hickory Street, Metuchen, NJ 08840 (US). YAO, Xiaorui; 95 Kelly Drive, Edison, NJ 08820 (US).  <b>(74) Agents:</b> GOLDMAN, Kenneth, M. et al.; Schering-Plough Corporation, Patent Dept., K-6-1 1990, 2000 Galloping Hill Road, Kenilworth, NJ 07033-0530 (US).		<b>(81) Designated States:</b> AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, NO, NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UZ, VN, YU, ZA, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  <b>Published</b> <i>With international search report.</i> <i>Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>
<b>(54) Title:</b> USE OF AZETIDINONE COMPOUNDS  <b>(57) Abstract</b>  <p>The use of azetidinone compounds that are inhibitors of cholesterol absorption as tools for discovering and characterizing proteins involved in trafficking or absorption of cholesterol and/or cholesteryl esters in biological systems is presented. These compounds can serve as tools for competitive binding assays to discover and characterize other chemical agents useful as cholesterol absorption inhibitors. New compounds of the present invention are also highly efficacious inhibitors of cholesterol absorption.</p>		

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## USE OF AZETIDINONE COMPOUNDS

### Field of the Invention

The present invention relates to the use of azetidinone compounds that are inhibitors of cholesterol absorption as tools for discovering and characterizing proteins involved in trafficking or absorption of cholesterol and/or cholesteryl esters in biological systems. Further, these compounds can serve as tools for competitive  
5 binding assays to discover and characterize other chemical agents useful as cholesterol absorption inhibitors. New compounds of the present invention are highly efficacious inhibitors of cholesterol absorption.

### Background of the Invention

10 Atherosclerotic coronary heart disease (CHD) represents the major cause for death and cardiovascular morbidity in the western world. Risk factors for CHD include hypertension, diabetes mellitus, family history, male gender, cigarette smoke and serum cholesterol. A total cholesterol level in excess of 225 to 250 mg/dl is associated with significant elevation of risk of CHD.

15 Cholesteryl esters are a major component of atherosclerotic lesions and the major storage form of cholesterol in arterial wall cells. Formation of cholesteryl esters is also a key step in the intestinal absorption of dietary cholesterol. Thus, inhibition of cholesteryl ester formation and reduction of serum cholesterol is likely to inhibit the progression of atherosclerotic lesion formation, decrease the accumulation of  
20 cholesteryl esters in the arterial wall, and block the intestinal absorption of dietary cholesterol.

The regulation of whole-body cholesterol homeostasis in humans and animals involves the regulation of dietary cholesterol and modulation of cholesterol biosynthesis, bile acid biosynthesis and the catabolism of the cholesterol-containing  
25 plasma lipoproteins. The liver is the major organ responsible for cholesterol biosynthesis and catabolism and for this reason, it is a prime determinant of plasma cholesterol levels. The liver is the site of synthesis and secretion of very low density

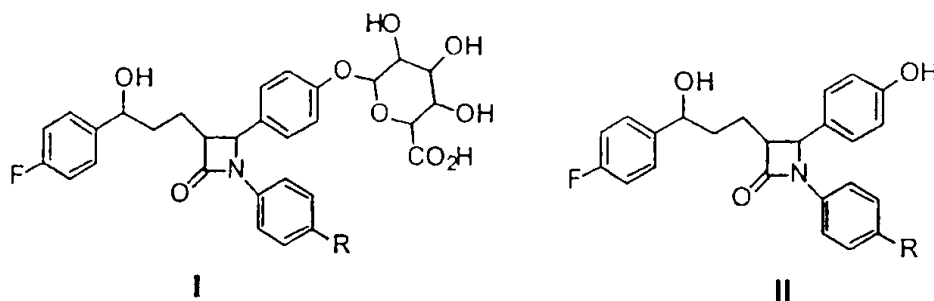
lipoproteins (VLDL) which are subsequently metabolized to low density lipoproteins (LDL) in the circulation. LDL are the predominant cholesterol-carrying lipoproteins in the plasma and an increase in their concentration is correlated with increased atherosclerosis.

5        When intestinal cholesterol absorption is reduced, by whatever means, less cholesterol is delivered to the liver. The consequence of this action is decreased hepatic lipoprotein production, and an increase in the hepatic clearance of plasma cholesterol, mostly as LDL. Thus, the net effect of inhibiting intestinal cholesterol absorption is a decrease in plasma cholesterol levels.

10        Certain azetidinone core structures have been reported to be useful in lowering cholesterol levels by decreasing intestinal cholesterol absorption. These related azetidinone cores and their synthesis are detailed in the following commonly assigned United States patents, the disclosures of which are incorporated, in their entirety, herein by reference: U.S. Patent Nos. 5,688,787; 5,698,548; 5,624,920; 5,631,365;  
15        5,633,246; 5,656,624; 5,744,467; and 5,767,115. The discovery of 2-azetidinones as potent and selective intestinal cholesterol absorption inhibitors has confirmed this mechanism as a key point of intervention for lowering cholesterol plasma levels and has validated the therapeutic value of such an approach.

20        The mechanism by which cholesterol moves from the lumen into the epithelial layer lining the small intestine is not well understood. Recent experimental evidence supports the notion of an active transport process mediated by a protein or proteins in the enterocyte brush border membrane rather than a simple diffusion model. Kinetic analysis and sterol specificity of cholesterol uptake as well as the structure-activity relationship studies of the cholesterol absorption inhibitors are consistent with a  
25        specific protein receptor/transporter regulated event. Potential molecules for this process have been proposed in recent years. However, the specific biochemical pathway responsible for cholesterol absorption remains to be defined.

Compounds I and II (especially II)



where R is fluorine, are potent inhibitors of cholesterol uptake in animal models and humans. The mechanism by which these compounds and related 2-azetidinones inhibit the uptake of cholesterol across the intestinal wall is not known. These compounds do not sequester bile acids or precipitate cholesterol. Nor do they potently inhibit HMG-CoA reductase, pancreatic lipase, or acyl-CoA cholesterol acyl transferase (ACAT). Understanding the mechanism by which these compounds inhibit cholesterol absorption will shed light on the biochemical pathways involved in the uptake of dietary and biliary cholesterol.

#### Summary of the Invention

In one aspect, this invention provides a method for identifying a protein involved in cholesterol absorption in a biological system, the method comprising the steps of: (a) providing a cDNA expression library capable of expressing a protein involved in cholesterol absorption in a biological system, said cDNA expression library comprising a plurality of cells capable of expressing different cDNAs; (b) screening said expression library by incubating cells from said library with a fluorescent cholesterol absorption inhibitor; (c) after step (b), identifying the cell or cells in said library that display the greatest amount of fluorescence; and (d) identifying the protein associated with the cDNA expressed by said cell or cells displaying the greatest amount of fluorescence.

In another aspect, this invention provides a method for assaying inhibitory agents for activity against cholesterol absorption, the method comprising the steps of:

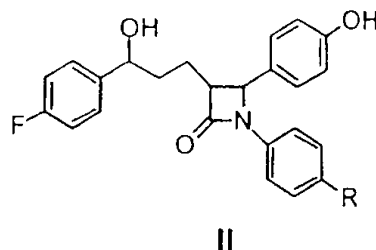
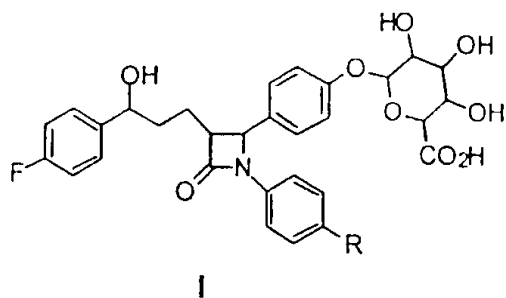
providing a cell capable of binding a fluorescent cholesterol absorption inhibitor; contacting said cell with a candidate inhibitory agent in the presence of said fluorescent cholesterol absorption inhibitor; and measuring the inhibition of the fluorescence of said cell.

5           In still another aspect, this invention provides a method for identifying inhibitory agents which inhibit the absorption of cholesterol into a cell membrane, said method comprising the steps of: (a) combining a fluorescent cholesterol absorption inhibitor, said cell membrane and a candidate inhibitory agent, under conditions wherein, but for the presence of said inhibitory agent, said fluorescent cholesterol absorption  
10 inhibitor is bound to the membrane; and (b) detecting the relative presence or absence of fluorescent cholesterol absorption inhibitor absorption bound to the membrane, wherein a relative absence of fluorescent cholesterol absorption inhibitor absorption indicates that said candidate inhibitory agent is an inhibitory agent which inhibits cholesterol absorption into the membrane.

15           In another aspect, this invention provides a method for identifying inhibitory agents which inhibit the absorption of cholesterol, said method comprising the steps of: (a) combining a labeled cholesterol absorption inhibitor, a cell expressing the scavenger receptor type B, class I (SR-BI) and a candidate inhibitory agent, under conditions wherein, but for the presence of said inhibitory agent, said labeled  
20 cholesterol absorption inhibitor binds to SR-BI; and (b) detecting the relative presence or absence of labeled cholesterol absorption inhibitor absorption bound to SR-BI, wherein a relative absence of labeled cholesterol absorption inhibitor absorption indicates that said candidate inhibitory agent is an inhibitory agent which inhibits SR-BI-mediated cellular cholesterol absorption.

25           In further aspects, this invention provides proteins and new inhibitory agents identified by the above methods. This invention also provides novel fluorescent cholesterol absorption inhibitors of formulas I and II:

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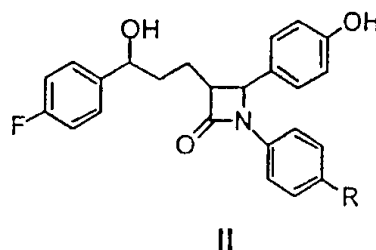
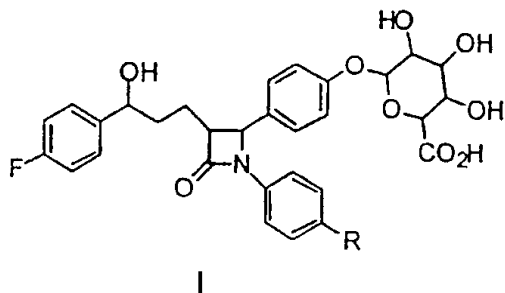


wherein R comprises a fluorescent moiety.

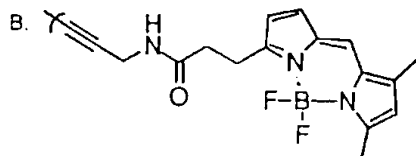
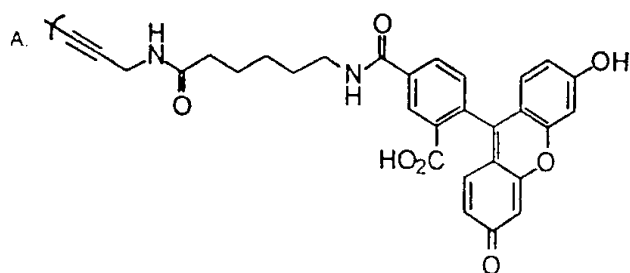
### Detailed Description of the Invention

#### 5 A. Novel fluorescent cholesterol absorption inhibitors

Novel fluorescent cholesterol absorption inhibitors of this invention include compounds of formulas I and II:

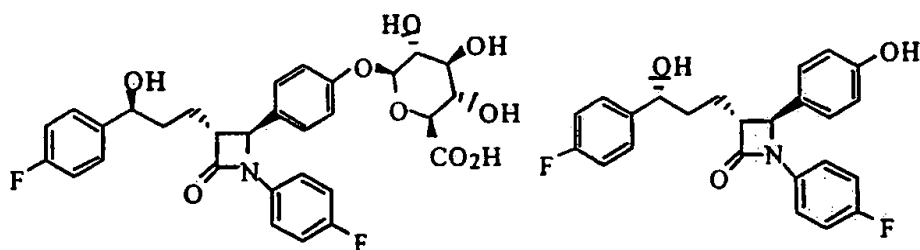


wherein R comprises a fluorescent moiety. In preferred embodiments, R is a fluorescent moiety linked by an alkynyl-containing tether group. In a particularly preferred embodiment, R is selected from the group consisting of:



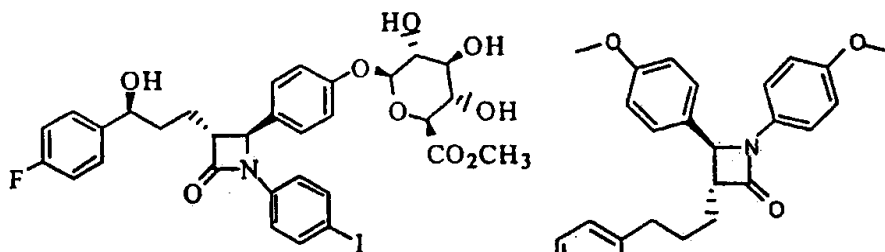
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Modification with fluorescent moieties A-B of the N-aryl ring of other azetidinone cholesterol absorption inhibitor core structures related to the above are contemplated as being within the scope of this invention as well. These related  
5 azetidinone cores and their synthesis are detailed in the following commonly assigned United States patents the disclosures of which are incorporated, in their entirety, herein by reference: U.S. Patent Nos. 5,688,787; 5,698,548; 5,624,920; 5,631,365; 5,633,246; 5,656,624; 5,744,467; and 5,767,115.  
Compounds in the following text are designated as:



IA

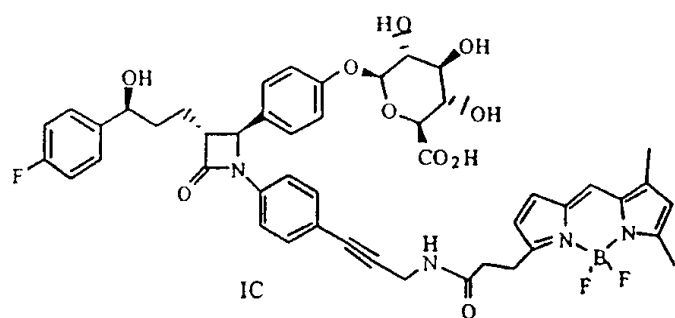
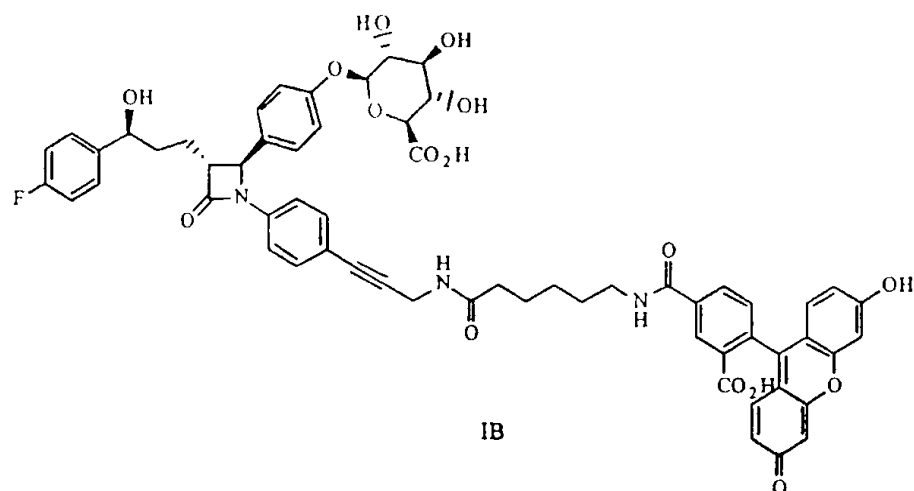
IIA



IIIA

IVA

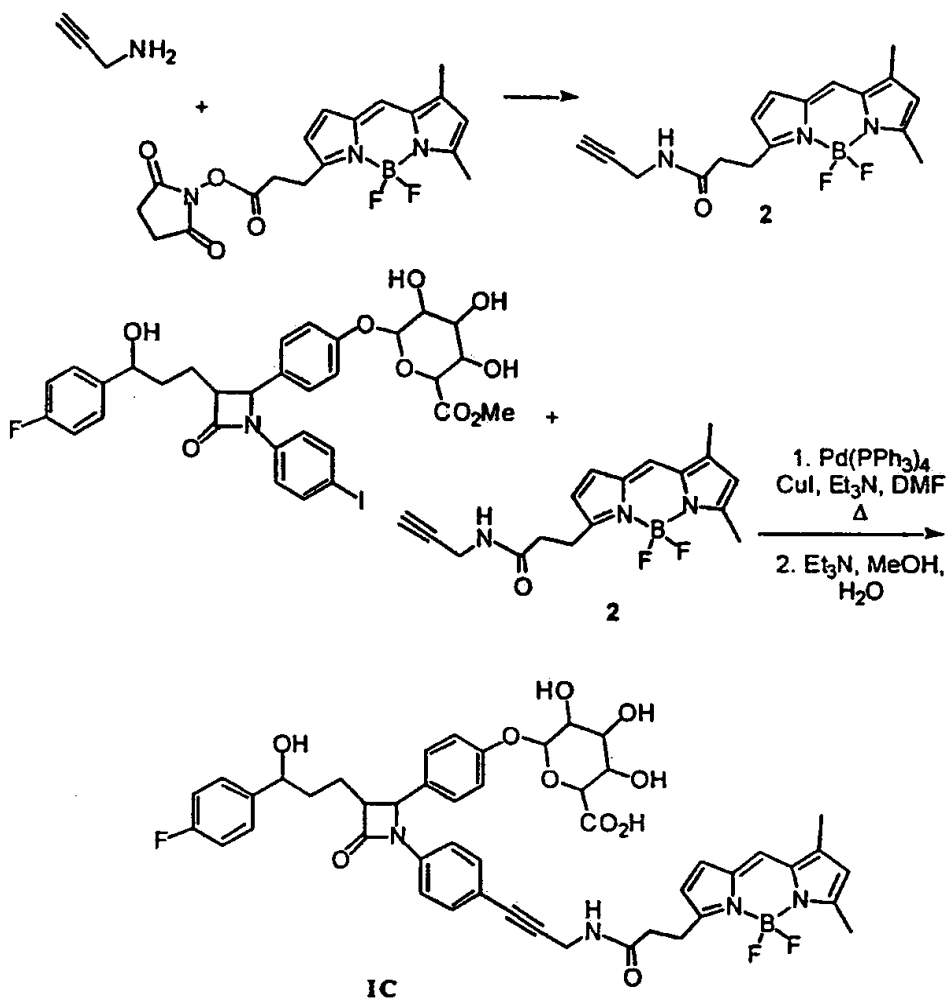




The synthetic route to these fluorescent azetidinones is shown in

5 Scheme 1. The fluorescent alkyne compounds are prepared from commercially available electrophilic fluorescent derivatives (see Handbook of Fluorescent Probes and Research Chemicals, Molecular Probes, Inc.) and propargyl amine. The syntheses of N-iodophenyl azetidinone derivatives have been described in previous patents, cited above. The fluorescent alkynylated material is then coupled to the N-  
10 iodophenylazetidinone via a palladium mediated coupling reaction. Deprotection of the glucuronide ester is accomplished as a final step in analogs bearing this moiety.

Scheme 1.



## 5 B. General Methods

The practice of the present invention generally employs conventional techniques of molecular biology, microbiology, recombinant DNA, and immunology, which are within the skill of the art. Such techniques are explained fully in the literature. See for example J. Sambrook et al, "Molecular Cloning; A Laboratory Manual (1989); "DNA Cloning", Vol. I and II (D. N. Glover ed. 1985); "Oligonucleotide

10 Synthesis" (M. J. Gait ed, 1984); "Nucleic Acid Hybridization" (B. D. Hames & S. J.

Higgins eds. 1984); "Transcription And Translation" (B. D. Hames & S. J. Higgins eds. 1984); "Animal Cell Culture" (R. I. Freshney ed. 1986); "Immobilized Cells And Enzymes" (IRL Press, 1986); B. Perbal, "A Practical Guide To Molecular Cloning" (1984); the series, "Methods In Enzymology" (Academic Press, Inc.); "Gene Transfer Vectors For Mammalian Cells" (J. H. Miller and M. P. Calos eds. 1987, Cold Spring Harbor Laboratory); Meth Enzymol (1987) 154 and 155 (Wu and Grossman, and Wu, eds., respectively); Mayer & Walker, eds. (1987), "Immunochemical Methods In Cell And Molecular Biology" (Academic Press, London); Scopes, "Protein Purification: Principles And Practice", 2nd Ed (Springer-Verlag, N.Y., 1987); and "Handbook Of Experimental Immunology", volumes I-IV (Weir and Blackwell, eds, 1986).

### C. Methods of screening cDNA libraries

cDNA libraries containing genes capable of expressing a protein involved in cholesterol absorption are prepared by standard techniques, such as described in the above references. Both prokaryotic and eukaryotic host cells are useful for expressing desired coding sequences when appropriate control sequences compatible with the designated host are used. Among prokaryotic hosts, *E. coli* is most frequently used. Expression control sequences for prokaryotes include promoters, optionally containing operator portions, and ribosome binding sites. Transfer vectors compatible with prokaryotic hosts are commonly derived from, for example, pBR322, a plasmid containing operons conferring ampicillin and tetracycline resistance, and the various pUC vectors, which also contain sequences conferring antibiotic resistance markers. These plasmids are commercially available. The markers may be used to obtain successful transformants by selection. Commonly used prokaryotic control sequences include the  $\beta$ -lactamase (penicillinase) and lactose promoter systems (Chang et al, Nature (1977) 198:1056), the tryptophan (trp) promoter system (Goeddel et al, Nuc Acids Res (1980) 8:4057) and the lambda-derived  $P_L$  promoter and N gene ribosome binding site (Shimatake et al, Nature (1981) 292:128) and the hybrid tac promoter (De Boer et al, Proc Nat Acad Sci USA (1983) 292:128) derived from sequences of the trp and lac UV5 promoters. The foregoing systems are particularly compatible with *E. coli*; if desired, other prokaryotic

hosts such as strains of *Bacillus* or *Pseudomonas* may be used, with corresponding control sequences.

Eukaryotic hosts include without limitation yeast and mammalian cells in culture systems. Yeast expression hosts include *Saccharomyces*, *Klebsiella*, *Picia*, and the like. *Saccharomyces cerevisiae* and *Saccharomyces carlsbergensis* and *K. lactis* are the most commonly used yeast hosts, and are convenient fungal hosts. Yeast-compatible vectors carry markers which permit selection of successful transformants by conferring prototrophy to auxotrophic mutants or resistance to heavy metals on wild-type strains. Yeast compatible vectors may employ the 2 $\mu$  origin of replication (Broach et al, *Meth Enzymol* (1983) 101:307), the combination of CEN3 and ARS1 or other means for assuring replication, such as sequences which will result in incorporation of an appropriate fragment into the host cell genome. Control sequences for yeast vectors are known in the art and include promoters for the synthesis of glycolytic enzymes (Hess et al, *J Adv Enzyme Reg* (1968) 7:149; Holland et al, *Biochem* (1978), 17:4900), including the promoter for 3-phosphoglycerate kinase (R. Hitzeman et al, *J Biol Chem* (1980) 255:2073). Terminators may also be included, such as those derived from the enolase gene (Holland, *J Biol Chem* (1981) 256:1385). Particularly useful control systems are those which comprise the glyceraldehyde-3 phosphate dehydrogenase (GAPDH) promoter or alcohol dehydrogenase (ADH) regulatable promoter, terminators also derived from GAPDH, and if secretion is desired, a leader sequence derived from yeast alpha-factor (see U.S. Pat. No. 4,870,008, incorporated herein by reference).

In addition, the transcriptional regulatory region and the transcriptional initiation region which are operably linked may be such that they are not naturally associated in the wild-type organism. These systems are described in detail in EPO 120,551, published October 3, 1984; EPO 116,201, published Aug. 22, 1984; and EPO 164,556, published Dec. 18, 1985, all of which are hereby incorporated herein by reference in full.

Mammalian cell lines available as hosts for expression are known in the art and include many immortalized cell lines available from the American Type Culture Collection (ATCC), including HeLa cells, Chinese hamster ovary (CHO) cells, baby

hamster kidney (BHK) cells, and a number of other cell lines. Suitable promoters for mammalian cells are also known in the art and include vital promoters such as that from Simian Virus 40 (SV40) (Fiers et al, Nature (1978) 273:113), Rous sarcoma virus (RSV), adenovirus (ADV), and bovine papilloma virus (BPV). Mammalian cells may also require terminator sequences and poly-A addition sequences. Enhancer sequences which increase expression may also be included, and sequences which promote amplification of the gene may also be desirable (for example methotrexate resistance genes). These sequences are known in the art.

Vectors suitable for replication in mammalian cells are known in the art, and may include vital replicons, or sequences which insure integration of the appropriate sequences into the host genome. For example, another vector used to express foreign DNA is Vaccinia virus. In this case the heterologous DNA is inserted into the Vaccinia genome. Techniques for the insertion of foreign DNA into the vaccinia virus genome are known in the art, and may utilize, for example, homologous recombination. The heterologous DNA is generally inserted into a gene which is non-essential to the virus, for example, the thymidine kinase gene (tk), which also provides a selectable marker. Plasmid vectors that greatly facilitate the construction of recombinant viruses have been described (see, for example, Mackett et al, J Virol (1984) 49:857; Chakrabarti et al, Mol Cell Biol (1985) 5:3403; Moss, in GENE TRANSFER VECTORS FOR MAMMALIAN CELLS (Miller and Calos, eds., Cold Spring Harbor Laboratory, N.Y., 1987), p. 10). Expression of the polypeptide then occurs in cells or animals which are infected with the live recombinant vaccinia virus.

Transformation may be by any known method for introducing polynucleotides into a host cell, including, for example packaging the polynucleotide in a virus and transducing a host cell with the virus, and by direct uptake of the polynucleotide. The transformation procedure used depends upon the host to be transformed. Bacterial transformation by direct uptake generally employs treatment with calcium or rubidium chloride (Cohen, Proc Nat Acad Sci USA (1972) 69:2110; T. Maniatis et al, "Molecular Cloning; A Laboratory Manual" (Cold Spring Harbor Press, Cold Spring Harbor, N.Y., 1982). Yeast transformation by direct uptake may be carried out using the method of Hinnen et al, Proc Nat Acad Sci USA (1978) 75:1929. Mammalian transformations by

direct uptake may be conducted using the calcium phosphate precipitation method of Graham and Van der Eb, *Virology* (1978) 52:546, or the various known modifications thereof. Other methods for introducing recombinant polynucleotides into cells, particularly into mammalian cells, include dextran-mediated transfection, calcium phosphate mediated transfection, polybrene mediated transfection, protoplast fusion, electroporation, encapsulation of the polynucleotide(s) in liposomes, and direct microinjection of the polynucleotides into nuclei.

Vector construction employs techniques which are known in the art. Site-specific DNA cleavage is performed by treating with suitable restriction enzymes under conditions which generally are specified by the manufacturer of these commercially available enzymes. In general, about 1 µg of plasmid or DNA sequence is cleaved by 1 unit of enzyme in about 20 µL buffer solution by incubation for 1-2 hr at 37°C. After incubation with the restriction enzyme, protein is removed by phenol/chloroform extraction and the DNA recovered by precipitation with ethanol. The cleaved fragments may be separated using polyacrylamide or agarose gel electrophoresis techniques, according to the general procedures described in *Meth Enzymol* (1980) 65:499-560.

Sticky-ended cleavage fragments may be blunt ended using *E. coli* DNA polymerase I (Klenow fragment) with the appropriate deoxynucleotide triphosphates (dNTPs) present in the mixture. Treatment with S1 nuclease may also be used, resulting in the hydrolysis of any single stranded DNA portions.

Ligations are carried out under standard buffer and temperature conditions using T4 DNA ligase and ATP; sticky end ligations require less ATP and less ligase than blunt end ligations. When vector fragments are used as part of a ligation mixture, the vector fragment is often treated with bacterial alkaline phosphatase (BAP) or calf intestinal alkaline phosphatase to remove the 5'-phosphate, thus preventing religation of the vector. Alternatively, restriction enzyme digestion of unwanted fragments can be used to prevent ligation. Ligation mixtures are transformed into suitable cloning hosts, such as *E. coli*, and successful transformants selected using the markers incorporated (e.g., antibiotic resistance), and screened for the correct construction.

Synthetic oligonucleotides may be prepared using an automated oligonucleotide synthesizer as described by Warner, DNA (1984) 3:401. If desired, the synthetic strands may be labeled with  $^{32}\text{P}$  by treatment with polynucleotide kinase in the presence of  $^{32}\text{P}$ -ATP under standard reaction conditions.

5 For routine vector constructions, ligation mixtures are transformed into *E. coli* strain HB101 or other suitable hosts, and successful transformants selected by antibiotic resistance or other markers. Plasmids from the transformants are then prepared according to the method of Clewell et al, Proc Nat Acad Sci USA (1969) 62:1159, usually following chloramphenicol amplification (Clewell, J Bacteriol (1972) 10 110:667). The DNA is isolated and analyzed, usually by restriction enzyme analysis and/or sequencing. Sequencing may be performed by the dideoxy method of Sanger et al, Proc Nat Acad Sci USA (1977) 74:5463, as further described by Messing et al, Nuc Acids Res (1981) 9:309, or by the method of Maxam et al, Meth Enzymol (1980) 65:499. Problems with band compression, which are sometimes observed in GC-rich 15 regions, were overcome by use of T-deazoguanosine according to Barr et al, Biotechniques (1986) 4:428.

Preferably, the cDNA libraries used in this invention are derived from the mRNA of cells or cell lines known to be involved in cholesterol absorption. In a preferred embodiment of this invention, the cDNA library is derived from the mRNA of 20 an intestinal epithelial cell. The cDNA libraries are then screened by incubating cells from the library with a fluorescent cholesterol absorption inhibitor, and identifying those cells that display the greatest amount of fluorescence, using, for example, fluorescence activated cell sorting (FACS). The cDNA expressed in that cell can then be identified using known techniques.

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#### D. Methods of Screening for Inhibitory Agents

New cholesterol absorption inhibitory agents are screened using methods of the invention. In general, a substrate is employed which mimics the protein's natural substrate, but which provides a quantifiable signal when bound. The signal is 30 preferably detectable by colorimetric or fluorometric means; however, other methods such as HPLC or silica gel chromatography, GC-MS, nuclear magnetic resonance,

and the like may also be useful. After optimum substrate and protein (or membrane or cell) concentrations are determined, a candidate inhibitory agent is added to the reaction mixture at a range of concentrations. The assay conditions ideally should resemble the conditions under which cholesterol absorption is to be inhibited in vivo, i.e., under physiologic pH, temperature, ionic strength, etc. Suitable inhibitors will exhibit absorption inhibition at concentrations which do not raise toxic side effects in the subject. Inhibitors which compete for binding may require concentrations equal to or greater than the substrate concentration, while inhibitors capable of binding irreversibly to the cholesterol binding site may be added in concentrations on the order of the protein concentration.

This invention provides several methods for assaying, identifying or screening inhibitory agents for activity against cholesterol absorption, including cellular techniques, membrane techniques, and techniques using proteins that are involved in cholesterol absorption, such as the one provided by the current invention, the scavenger receptor type B, class I (SR-BI). SR-BI was first described by Acton et al., *J. Biol. Chem.* (1994) 269:21003-21009. Screening for inhibitory agents is accomplished as described below.

In the cellular method, cells capable of binding a fluorescent cholesterol absorption inhibitor are used. In a preferred embodiment, the cells are involved in cholesterol absorption in a biological system. The cells are contacted with candidate inhibitory agents in the presence of the fluorescent cholesterol absorption inhibitor; the inhibition of the fluorescence of said cells are measured.

In the membrane method, a membrane capable of absorbing cholesterol is used. In a preferred embodiment of this invention, the membranes are from the intestinal brush border. The membranes are contacted with candidate inhibitory agents in the presence of the fluorescent cholesterol absorption inhibitor; and the relative presence or absence of fluorescent cholesterol absorption inhibitor absorption bound to the membrane is measured. A relative absence of fluorescent cholesterol absorption inhibitor absorption indicates that said candidate inhibitory agent is an inhibitory agent which inhibits cholesterol absorption into the membrane.



In the protein method, a cell expressing the scavenger receptor type B, class I (SR-BI) is used. The cell is contacted with a labeled cholesterol absorption inhibitor and a candidate inhibitory agent under conditions wherein, but for the presence of said inhibitory agent, the labeled cholesterol absorption inhibitor binds to SR-BI, and  
5 the relative presence or absence of labeled cholesterol absorption inhibitor absorption bound to SR-BI is detected. A relative absence of labeled cholesterol absorption inhibitor absorption indicates that the candidate inhibitory agent is an inhibitory agent which inhibits SR-BI-mediated cellular cholesterol absorption.

The methods of assaying or selecting inhibitory agents of this invention use  
10 labeled cholesterol absorption inhibitors. The inhibitors themselves are described in section A, above. The labels used in the assays of invention can be primary labels (where the label comprises an element which is detected directly) or secondary labels (where the detected label binds to a primary label, e.g., as is common in immunological labeling). An introduction to labels, labeling procedures and detection  
15 of labels is found in Polak and Van Noorden (1997) Introduction to Immunocytochemistry, second edition, Springer Verlag, N.Y. and in Haugland (1996) Handbook of Fluorescent Probes and Research Chemicals, a combined handbook and catalogue Published by Molecular Probes, Inc., Eugene, OR. Primary and secondary labels can include undetected elements as well as detected elements.  
20 Useful primary and secondary labels in the present invention can include spectral labels, which include fluorescent labels such as fluorescent dyes (e.g., fluorescein and derivatives such as fluorescein isothiocyanate (FITC) and Oregon Green™, rhodamine and derivatives (e.g., Texas red, tetramethylrhodamine isothiocyanate (TRITC), etc.), digoxigenin, biotin, phycoerythrin, AMCA, CyDyes™, and the like),  
25 radiolabels (e.g., <sup>3</sup>H, <sup>125</sup>I, <sup>35</sup>S, <sup>14</sup>C, <sup>32</sup>P, <sup>33</sup>P, etc.), enzymes (e.g., horseradish peroxidase, alkaline phosphatase etc.) spectral colorimetric labels such as colloidal gold or colored glass or plastic (e.g. polystyrene, polypropylene, latex, etc.) beads. The label may be coupled directly or indirectly to the cholesterol absorption inhibitor according to methods well known in the art. As indicated above, a wide variety of  
30 labels may be used, with the choice of label depending on sensitivity required, ease of conjugation with the compound, stability requirements, available instrumentation,

and disposal provisions. In general, a detector which monitors a protein/inhibitory agent interaction is adapted to the particular label which is used. Typical detectors include spectrophotometers, phototubes and photodiodes, microscopes, scintillation counters, cameras, film and the like, as well as combinations thereof. Examples of  
5 suitable detectors are widely available from a variety of commercial sources known to persons of skill.

Preferred labels include those which utilize 1) chemiluminescence (using horseradish peroxidase or alkaline phosphatase with substrates that produce photons as breakdown products) with kits being available, e.g., from Molecular Probes,  
10 Amersham, Boehringer-Mannheim, and Life Technologies/Gibco BRL; 2) color production (using both horseradish peroxidase or alkaline phosphatase with substrates that produce a colored precipitate) (kits available from Life Technologies/Gibco BRL, and Boehringer-Mannheim); 3) fluorescence (e.g., using Cy-5 (Amersham), fluorescein, and other fluorescent tags); 5) radioactivity. Other  
15 methods for labeling and detection will be readily apparent to one skilled in the art.

Fluorescent labels are highly preferred labels, having the advantage of requiring fewer precautions in handling, and being amenable to high-throughput visualization techniques (optical analysis including digitization of the image for analysis in an integrated system comprising a computer). Preferred labels are  
20 typically characterized by one or more of the following: high sensitivity, high stability, low background, low environmental sensitivity and high specificity in labeling. Fluorescent moieties, which are incorporated into the labels of the invention, are generally known, including Texas red, digoxigenin, biotin, 1- and 2-aminonaphthalene, p,p'-diaminostilbenes, pyrenes, quaternary phenanthridine salts,  
25 9-aminoacridines, p,p'-diaminobenzophenone imines, anthracenes, oxacarbocyanine, merocyanine, 3-aminoequilenin, perylene, bis-benzoxazole, bis-p-oxazolyl benzene, 1,2-benzophenazin, retinol, bis-3-aminopyridinium salts, hellebrigenin, tetracycline, sterophenol, benzimidazolylphenylamine, 2-oxo-3-chromen, indole, xanthen, 7-hydroxycoumarin, phenoxazine, caliculate, strophanthidin, porphyrins,  
30 triarylmethanes, flavin and many others. Many fluorescent tags are commercially available from the SIGMA chemical company (Saint Louis, Mo.), Molecular Probes,

R&D systems (Minneapolis, Minn.), Pharmacia LKB Biotechnology (Piscataway, N.J.), CLONTECH Laboratories, Inc. (Palo Alto, Calif.), Chem Genes Corp., Aldrich Chemical Company (Milwaukee, Wis.), Glen Research, Inc., GIBCO BRL Life Technologies, Inc. (Gaithersburg, Md.), Fluka ChemicaBiochemika Analytika (Fluka Chemie AG, Buchs, Switzerland), and Applied Biosystems (Foster City, Calif.), as well as many other commercial sources known to one of skill.

The labels are covalently bound to the novel cholesterol inhibitors of the invention by a tether group. The tether group can be any moiety capable of covalently linking to the inhibitors and to the labels. Preferred groups are substituted or unsubstituted alkylene, alkenylene or alkynylene of 1 to 10 carbon atoms, more preferably 1 to 4 carbon atoms. Particularly preferred groups are unsubstituted alkynylenes.

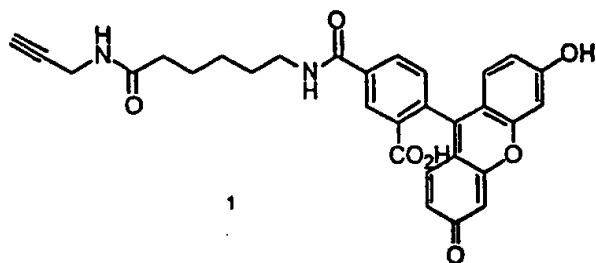
The candidate and actual inhibitory agents of this invention will preferably take the form of organic compounds, particularly compounds which inhibit the absorption of cholesterol mediated by SR-BI and other proteins<sup>3</sup>. Other inhibitory agents may be proteins, particularly antibodies and antibody derivatives. Recombinant expression systems may be used to generate quantities of SR-BI sufficient for production of monoclonal antibodies (MAbs) specific for SR-BI. Suitable antibodies for cholesterol absorption inhibition will bind to SR-BI in a manner reducing or eliminating the cholesterol binding activity, typically by obscuring the binding site. Suitable MAbs may be used to generate derivatives, such as Fab fragments, chimeric antibodies, altered antibodies, univalent antibodies, and single domain antibodies, using methods known in the art.

## EXAMPLES

### EXAMPLE 1

A. PREPARATION OF 6-(6-HYDROXY-3-OXO-3H-XANTHEN-9-YL)-N-(5-PROP-2-YNYLCARBAMOYL-PENTYL)-ISOPHTHALAMIC ACID (1):

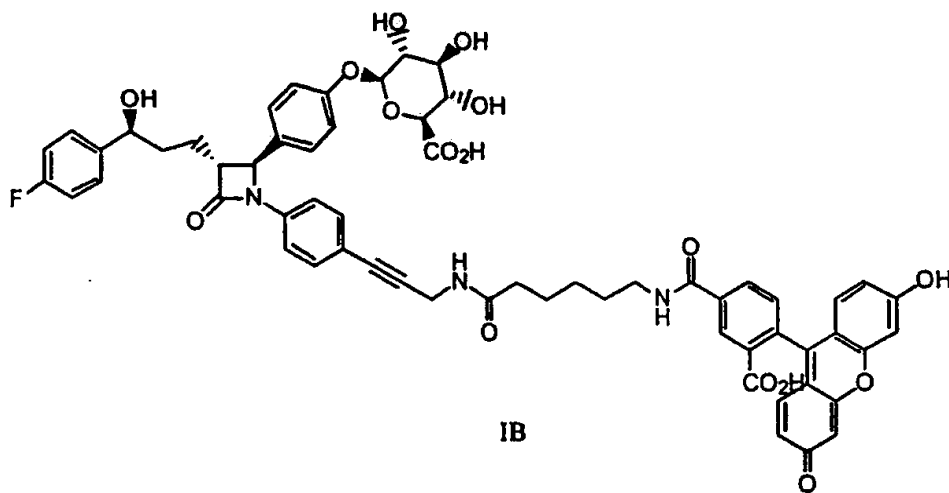
18



1

Added propargyl amine (5 $\mu$ L, ~4mg, 73 $\mu$ mol) to 0.3mL DMF containing 3-5  
 drops 0.2M aqueous NaHCO<sub>3</sub>, then added 6-(Fluorescein-5-carboxamido)hexanoic  
 acid, succinimidyl ester (5-SFX; Molecular Probes, Inc.) (5 mg, 8.5 $\mu$ mol) to the  
 mixture. Stirred the bright orange solution for 3h at room temperature. Concentrated  
 in vacuo and chromatographed the residue over ~5g of silica gel eluting with  
 10%MeOH in dichloromethane to give the desired terminal alkyne (1) as a yellow  
 brown fluorescent oil. calcd m/z for C<sub>30</sub>H<sub>27</sub>N<sub>2</sub>O<sub>7</sub> = 527; found m/z = 527.

B. PREPARATION OF 1-O-[4-[TRANS-(3R,4S)-1-[4-[3-[[6-[[3-CARBOXY-4-(6-  
 HYDROXY-3-OXO-3H-XANTHEN-9-YL)PHENYL]CARBONYL]AMINO]-1-  
 OXOHXYL]AMINO]-1-PROPENYL]PHENYL]-3-[3-(S)-HYDROXY-3-(4-  
 FLUOROPHENYL)PROPYL]-2-OXO-4-AZETIDINYL]PHENYL]-BETA-D-  
 GLUCORONIC ACID (IB):



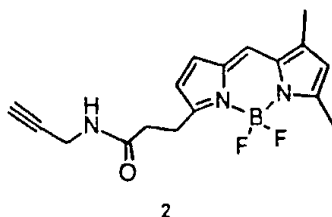
IB

To the iodophenylazetidinone glucuronide methyl ester IIIA (disclosed in Patent Number 5756,470, May 26, 1998) (6.7mg, 9.47 $\mu$ mol) in 0.8mL DMF, was added the alkyne (1) (14.8mg, 28 $\mu$ mol). Bubbled argon through the reaction mixture, then  
5 added palladium tetrakis(triphenylphosphine) (1.4mg, 1.2 $\mu$ mol), copper(I) iodide (0.8mg, 4.2 $\mu$ mol), and triethylamine (8 $\mu$ L, 57 $\mu$ mol). Stirred the reaction mixture overnight under argon. Concentrated the reaction onto a prep TLC plate and eluted with 25% MeOH in dichloromethane. Collected the major fluorescent band and purified further by reverse phase HPLC over a C-18 column eluting with 10-100%  
10 acetonitrile in water affording the desired fluorescent methyl ester as an orange solid. FABMS: calcd m/z for C<sub>61</sub>H<sub>57</sub>N<sub>3</sub>FO<sub>16</sub> (M+1)<sup>+</sup> = 1106.4; found m/z = 1106.5.

The glucuronide methyl ester was then hydrolyzed by dissolving 9mg of the compound in 2mL of a 7:2:1 mixture of water:triethylamine: methanol and stirring for 1h. Evaporation to dryness gave the desired fluorescent azetidinone glucuronide,  
15 (IB) as an orange solid. ES-MS: calcd m/z for C<sub>60</sub>H<sub>55</sub>N<sub>3</sub>FO<sub>16</sub> (M+1)<sup>+</sup> = 1092.3; found m/z = 1092.3.

## EXAMPLE 2

20 A. PREPARATION OF 4,4-DIFLUORO-5,7-DIMETHYL-4-BORA-3a,4a-DIAZA-s-INDACENE-3-N-PROP-2-YNYL-PROPIONAMIDE (2):



To propargyl amine (8 $\mu$ L, ~6.4mg, ~117 $\mu$ mol) in 0.5mL DMF and 1 drop of  
25 0.1M NaHCO<sub>3</sub>, was added 4,4-difluoro-5,7-dimethyl-4-bora-3a,4a-diaza-s-indacene-3-propionic acid, succinimide ester (BODIPY FL, SE; Molecular Probes, Inc.) (10mg, 25.7 $\mu$ mol) in 0.1mL DMF. Stirred the reaction at room temperature for 12h. Concentrated the mixture in vacuo and chromatographed the residue over ~5g silica

gel eluting with 1% MeOH in dichloromethane to give the desired alkene (2) as a